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AMENDMENTS TO THE CLAIMS

Please amend the claims as follows:

- 1. (Currently Amended) A vaccine for the treatment of <u>a</u> disease caused by <u>a</u> pathogenic Neisseria, the vaccine comprising <u>an at least one immunogenic component, said immunogenic component being an epitope on a Neisseria lipopolysaccharide inner core characterized by the presence of a phosphoethanolamine moiety linked to a 3, 6, 7, or a combination thereof, position of HepII of the inner core, based on the inner core of a Neisseria lipopolysaccharide, LPS, wherein said vaccine and being is capable of eliciting functional protective and/or immunoprophylactic antibodies against a majority of the strains within the species of the pathogenic Neisseria pathogenic Neisseria strain.</u>
- 2. (Currently Amended) A vaccine according to claim The vaccine of claim 1, wherein the said immunogenic component is capable of eliciting functional protective and/or immunoprophylactic antibodies against at least 60% of the strains within the species of the pathogenic Neisseria pathogenic Neisseria strains.
- 3. (Currently Amended) A vaccine according to claim The vaccine of claim 2, wherein the said immunogenic component is capable of eliciting functional protective and/or immunoprophylactic antibodies against at least 70% of the strains within the species of the pathogenic Neisseria pathogenic Neisseria strains.
- 4. (Currently Amended) A vaccine according to claim The vaccine of claim 1, wherein the immunogenic component is substantially free from outer core lipopolysaccharide.
- 5. (Currently Amended) A vaccine according to claim The vaccine of claim 1, wherein the species of the pathogenic Neisseria is Neisseria meningitidis.
- 6. (Currently Amended) A vaccine-according to claim The vaccine of claim 5, wherein the antibodies are elicited by and specifically recognize an [[the]] immunogenic component of [[in]] at least 50 % of group B strains of Neisseria meningitidis.
- 7. (Currently Amended) A vaccine according to claim The vaccine of claim 5, wherein the antibodies are elicited by and specifically recognize an [[the]] immunogenic component of [[in]] at least 60% of group B strains of Neisseria meningitidis.
- 8. (Currently Amended) A vaccine-according to claim The vaccine of claim 5, wherein the antibodies are elicited by and specifically recognize an [[the]] immunogenic component of [[in]] at least 70% of group B strains of Neisseria meningitidis.

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9. (Currently Amended) A vaccine according to claim The vaccine of claim 1, wherein the immunogenic component comprises [[of]] or consists of an epitope which is a part or all of the inner core structure of a of said Neisseria LPS inner core, is derived purified from this inner core, is a synthetic version of the inner core, or is a functional structural equivalent thereof.

- 10. (Currently Amended) A vaccine according to elaim The vaccine of claim 1, wherein the immunogenic component is an epitope on the LPS inner core characterized by the presence of a phosphoethanolamine moiety linked to the 3-position at HepII of the inner core, or is a functional structural equivalent thereof.
- 11. (Currently Amended) A vaccine according to elaim The vaccine of claim 1, wherein said immunogenic component is an epitope on the LPS inner core which comprises a glucose residue at HepI.
- 12. (Currently Amended) A vaccine according to claim The vaccine of claim 1 or 2, wherein the immunogenic component is an epitope on the LPS inner core which comprises an N-acetyl glucosamine at HepII of the inner core LPS.
- 13. (Currently Amended) A vaccine according to claim The vaccine of claim 1, wherein the inner core LPS consists of an inner core oligosaccharide attached to lipid A, with the general formula as shown:

Kdo
$$\alpha - (2,4)$$

$$Glc - \beta - (1,4) - HepI - \alpha - (1,5) - Kdo - \alpha - (2,6) - Lipid A$$

$$\alpha - (1,3)$$

$$R2$$

$$R1 - 3 - HepII$$

$$\alpha - (1,2)$$

$$R3$$

$$R4 - GlcNAc$$

where R1 is a substituent at the 3-position of HepII, and is hydrogen or Glc- α -(1, or phosphoethanolamine; R2 is a substituent at the 6-position of HepII, and is hydrogen or phosphoethanolamine; R3 is a substituent at the 7-position of HepII, and is hydrogen or phosphoethanolamine, and R4 is acetyl or hydrogen at the 3-position, 4-position or 6-position of the G1cNAc residue, or any combination thereof; and where Glc is D-glucopyranose; Kdo is 3-deoxy-D-manno-2-octulosonic acid; Hep is L-glycero-D-

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manno-heptose, and G1cNAc is 2-acetamido-2-deoxy-D-glucopyranose.

- 14. (Currently Amended) A vaccine according to claim-1, A vaccine for the treatment of a disease caused by a pathogenic Neisseria, the vaccine comprising at least one immunogenic component wherein said immunogenic component is reactive with the B5 a monoclonal antibody produced by the hybridoma deposited under accession number IDAC 260900-1, wherein said vaccine is capable of cliciting protective and/or immunoprophylactic antibodies against a pathogenic Neisseria strain.
- 15. (Currently Amended) The vaccine of claim 1, A vaccine comprising a few immunogenic component[[s]] based on the inner core of a Neisseria lipopolysaccharide, LPS, and being capable of cliciting functional antibodies against a majority of the strains within the species of the pathogenic Neisseria, further comprising a second immunogenic component, said second immunogenic component being an epitope on a Neisseria lipopolysaccharide inner core characterized by the presence of a phosphoethanolamine moiety linked to a 3, 6, 7, or a combination thereof, position of HepII of the inner core, wherein said phosphoethanolamine moiety of said second immunogenic component is linked to a different position of said HepII of the inner core than said phosphoethanolamine moiety of the immunogenic component of claim 1.
- 16. (Currently Amended) A vaccine according to claim 15 The vaccine of claim 1, and including an immunogenic component as defined in any of claims 1 to 14 wherein said immunogenic component has two phosphoethanolamine moieties located at two positions on said HepII, a first said position being the 3- position, and a second said position being the 6- or the 7- position.
- 17. (Currently Amended) A vaccine according to claim The vaccine of claim 15, wherein the said few immunogenic components vaccine elicits functional protective and/or immunoprophylactic antibodies against [[in]] at least 85% of the strains within the species of the pathogenic Neisseria pathogenic Neisseria strains.
- 18. (Currently Amended) A vaccine according to claim The vaccine of claim 17, wherein the said few immunogenic components vaccine elicits functional protective and/or immunoprophylactic antibodies against [[in]] at least 95% of the strains within the species of the pathogenic Neisseria pathogenic Neisseria strains.
- 19. Cancelled.
- 20. (Currently Amended) A vaccine according to claim The vaccine of claim 1, wherein [[the]] said immunogenic component element of the vaccine is an epitope accessible on [[the]] a bacterium in the presence of a lipopolysaccharide outer core of a bacterial capsule.

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21. (Currently Amended) A vaccine-according to claim The vaccine of claim 1, wherein said immunogenic element is comprising one or more immunogen components which are capable of stimulating antibodies which are opsonic for pathogenic Neisseria.

- 22. (Currently Amended) A vaccine according to claim The vaccine of claim 1 for the treatment of a condition characterized by Neisseria meningitidis infection.
- 23. (Currently Amended) A vaccine according to claim The vaccine of claim 22 for the treatment of a condition characterized by Neisseria meningitidis group B infection.
- 24. (Currently Amended) A vaccine according to Use of the vaccine of claim 1, for the treatment of meningitis, septicaemia or pneumonia or any other manifestation of systemic or local disease occasioned by a Neisseria meningitidis strain that is specifically recognized by antibodies elicted by said vaccine.
- 25. (Currently Amended) <u>Use of the vaccine of A vaccine according to claim 1</u> for the <u>prevention-treatment</u> of urethritis, salpingitis, cervicitis, proctitis, pharyngitis, pelvic inflammatory disease or <u>any</u> other manifestation of systemic or local disease occasioned by <u>a Neisseria gonorrhoeae strain that is specifically recognized by antibodies elicted by said vaccine.</u>
- 26. (Currently Amended) A vaccine according to claim The vaccine of claim 1, which is a conjugated vaccine.
- 27. (Currently Amended) A vaccine according to claim The vaccine of claim 1, which is derived from a wherein said Neisseria lipopolysaccharide inner core is a commensal Neisseria inner core.
- 28. (Currently Amended) A vaccine according to claim The vaccine of claim 27, wherein the commensal Neisseria is Neisseria lactamica.
- 29-41. Cancelled.
- 42. (New) A vaccine for the treatment of disease caused by pathogenic *Neisseria*, comprising a *galE* mutant *Neisseria* bacterium, wherein said vaccine elicits protective and/or immunoprophylactic antibodies against a majority of pathogenic Neisseria strains.
- 43. (New) The vaccine of claim 42, wherein said Neisseria bacterium is N. meningitidis.
- 44. (New) The vaccine of claim 42, wherein said N. meningitidis is a group B N. meningitidis.
- 45. (New) The vaccine of claim 42, wherein said *N. meningitidis* bacterium comprises a lipopolysaccharide inner core characterized by the presence of a phosphoethanolamine moiety linked to a 3, 6, 7, or a combination thereof, position of HepII of the inner core.

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46. (New) The vaccine of claim 42, wherein said N. meningitidis bacterium is killed or inactivated.

47. (New) The vaccine of claim 42, wherein said N. meningitidis bacterium is attenuated.